TIMING OF LATE PALAEOPROTEROZOIC METAMORPHISM IN THE NORTHERN BELOMORIAN BELT, WHITE SEA REGION: CONCLUSIONS FROM U-Pb ISOTOPIC DATA AND P-T EVIDENCE

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Brilliant-looking zircon crystals from a garnet-amphibolitic, metamorphosed mafic dyke at Lyagkomina in the Lake Kovdozero area, northern Belomorian Belt, most probably mark the 1875±4 Ma age of peak late Palaeoproterozoic regional metamorphism which followed upon dyke intrusion associated with deformation of the Lapland-Kola orogenic tectonic collage. U-Pb ages of titanites (ca. 1860 Ma) and rutiles (ca. 1750 Ma) help reconstruct the retrograde path and the cooling rates which decreased with time. The new titanite and rutile ages agree well with similar age data previously obtained on a regional scale from the central part of the belt and suggest absence of variation in mineral ages as a function of lithological differences. P-T analysis of the metamorphism and jump uplift with unroofing of ca. 9-10 km of the crust in the Belomorian Belt previously thickened by late Palaeoproterozoic orogenic thrusting and stacking.

Key words: metamorphism, P-T conditions, absolute age, U/Pb, zircon, titanite, rutile, Paleoproterozoic, Lyagkomina, Russian Federation

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INTRODUCTION

The Belomorian Belt along the south-western shore of the White Sea (Fig. 1) is one of the principal crustal units in the Archaean, eastern part of the Baltic (Fennoscandian) Shield. It is structurally complex and has undergone repeated metamorphism, the different mineral associations overprinting each other. Largely because of this complexity, the Belomorian Belt has long been regarded as the ancient core of the Archaean domain.

A completely different interpretation, however, was proposed by Gaál and Gorbatschev in 1987. These authors see the Belomorian as a mobile belt which in the Neoarchaean had collided with the Karelian protocraton and in late Palaeoproterozoic times had been compressed between that protocraton and the Archaean terranes of the Kola Peninsula. Subsequent work confirmed the validity of this reinterpretation. The present study focuses on the late Palaeoproterozoic development of the Belomorian Belt. The U-Pb age of brilliant-looking zircons in a deformed and metamorphosed mafic dyke is assumed to date approximately the peak of late Palaeoproterozoic metamorphism, and provide a reference point which cannot be obtained from the surrounding high-grade Archaean country rocks where zircons are complex and separate grains of relevant age could not be detected. The dated dyke also makes it possible to distinguish between early and late stages of deformation and metamorphism.

In addition, U-Pb dating of titanite and rutile opens a way to assess the final stages of the late Palaeoproterozoic metamorphic evolution. In that regard there is a close relationship between the present paper and the titanite – rutile datings recently published by Bibikova et al. (2001). However, while the latter study aimed at regional, large-scale investigation of the Belomorian Belt



Fig. 1. Sketch map of the Kovdozero-Tolstik area in the northern part of the Belomorian Belt (modified after Balagansky et al. 1986, Stepanov & Slabunov 1989, Miller & Milkevich 1995). The Lyagkomina Dam area (Fig 2) is shown as a rectangle.

and its south-western, Karelian foreland, the present project concentrates on an area of limited size (Fig. 1), where a range of lithologies can be compared.

REGIONAL SETTING: EVOLUTION OF THE BELOMORIAN BELT

The Belomorian Belt was formed in the Neoarchaean after ca. 2.9 Ga, apparently outboard the Karelian protocraton to which it was accreted ca. 2.7 Ga ago (Bibikova et al. 1993 and 1996, Bogdanova & Bibikova 1993, Miller & Mil'kevich 1995). The accretion is considered to have involved subduction beneath the protocraton as well as thrusting of the Belomorian rocks to the west. The one-time thrust planes, like the one along Lake Seryak (Fig. 1), are still among the most conspicuous mega-features of the Belomorian Belt.

The lithologies formed during the early development stages of the belt comprise the Chupa and Khetolambina rocks, terms which originally were used in a stratigraphic sense. The Chupa rocks are aluminous gneisses with greywacke and pelite protoliths. The Khetolambina rocks, in contrast, are principally basaltic to andesitic amphibolites. Both these supracrustal lithologies have been intruded by plutonic rocks of the TTG (tonalite-trondhjemite-granodiorite) association. Neoarchaean metamorphism featured a complicated sequence of events but, in brief, it involved a granulitic phase at moderately high pressures, succeeded by retrogression until ca. 2.6 Ga.

In the late Palaeoproterozoic, the Belomorian Belt was once more involved in orogenic processes, now during the Lapland-Kola orogeny at ca. 1.9-1.8 Ga (e.g. Bridgwater et al. 1992). That orogeny included continent collision between the Archaean terranes of the Kola Peninsula and the combined Karelian-Belomorian protocraton which had been developed in the Neoarchaean. The Belomorian Belt became involved in thrusting, was thickened, and finally subjected to folding and regional metamorphism. Its Neoarchaean suture boundary towards the Karelian crustal domain continued to act as a discontinuity and therefore became the site of particularly intense and long-lived hydrothermal and pegmatite-generating igneous activity with the youngest U-Pb titanite ages in the whole Belomorian region (Bibikova et al. 2001).

In the present context, it is of interest to note that during the late Palaeoproterozoic the titanite ages were completely reset and new rutile crystallized in the Belomorian Belt but not in the former cratonic nucleus in Karelia (Bibikova et al. 2001). In our recent U-Pb zircon studies of the Belomorian gneisses we have found a number of cases where discordant U-Pb zircon data point at lower concordia-discordia intercepts of roughly 1.9-1.8 Ga (Bibikova et al. 1996). On the other hand, zircon newly formed during that time span appears to be rare, and independent new crystals are few if present at all.

The time between the two orogenies comprised an early stage of rifting. Then followed bimodal plutonic igneous activity with a dominant mafic mode and associated 2.45 - 2.35 Ga high- to medium-grade metamorphism together with shearing and folding in the lower crust (Bogdanova & Bibikova 1993; Bridgwater et al. 1994, Bogdanova 1996, and references in these articles).

GEOLOGY OF THE LYAGKOMINA AND TUPAYA BAY LOCALITIES

All the three principal Neoarchaean lithologies of the Belomorian Belt are exposed in the Lyagkomina Dam area by Lake Kovdozero (Fig. 2). In the north-west, almost monomineralic amphibolites associate with diopside-clinozoisite- and garnetdominated gneisses which belong to the Khetolambina complex of ultramafic and mafic metamorphosed volcanics and intrusions. In the southeast, garnet-biotite-kyanite-muscovite-bearing Chupa gneisses border on foliated TTG-type plutonic rocks.

Three main stages of deformation can be distinguished in the area (Fig. 2). In the Neoarchaean, ductile overthrusting of Khetolambina amphibolites and mafic gneisses over the Chupa-type aluminous gneisses and some small-scale recumbent folding occurred. Thereafter followed an as-



Fig. 2. Geological map of the Lyagkomina Dam locality, based on mapping by S.B. in 1987. Points 91011– 91014 indicate localities sampled for isotope analysis.

sumedly early Palaeoproterozoic stage when the Neoarchaean supracrustal rocks together with the TTG-type intrusions were deformed into major, moderately inclined, close folds with NW-trending axes. We associate this stage of folding with the one that appears to be related to shearing, highgrade metamorphism and bimodal gabbroic-granitic igneous activity at 2.5-2.4 Ga in the Tolstik area by the White Sea (Fig. 1, cf. Bogdanova & Bibikova 1993, Bridgwater et al. 1994, Bogdanova 1996, Lobach-Zhuchenko et al. 1998, Claesson et al. in preparation).

The metamorphosed dykes, one of which has been investigated and dated in the present study (locality 91014 in Fig. 2), appear to separate that stage of deformation from a late Palaeoproterozoic one at a time when preexisting folds were flattened and folding of the dyke into small recumbent folds occurred. Subsequently followed boudinaging as well as migmatitic and pegmatitic veining in the enclosing aluminous gneisses. The dyke itself, however, is only weakly affected along its contacts with the host rocks.

Because of suspected disequilibrium among the

minerals of the metamorphic assemblages, it has been diffucult to assign specific P-T values to the late Neoarchaean evolution. We estimate, however, that the garnet \pm kyanite \pm biotite + plagioclase + quartz assemblage in the Chupa gneisses (Bibikova et al. 1993) had reached temperatures of 750-800°C at pressures of 10-12 kbar during the peak phase. A plausible hypothesis is that this occurred in association with the intrusion of ca. 2.73-Ga diorites along thrusts. Later, a retrograde amphibolitic stage was completed presumably by ca. 2.6 Ga. However, similar metamorphism has also been encountered among 2.45-2.40 Ga mafic intrusions in the Tolstik Peninsula (Fig. 1, cf. Bogdanova 1996 and references therein). A final phase of low-grade retrogression took place during uplift (cf. Miller et al. 1995, Skiöld et al. 1997) from about 1.9 Ga ago.

The Tupaya Bay locality (SE of Lyagkomina in Fig. 1) allows a relatively better documentation of several stages of Neoarchaean metamorphism which here reached conditions of 800°C at 10 kbar (cf. Balagansky et al. 1986, Bogdanova & Bibikova 1993, Lobach-Zhuchenko et al. 1993). The 2.45-Ga granitic intrusions of this locality were also deformed, and this late development of amphibolite-facies mineral assemblages tentatively corresponds to the stage of dyke intrusions and subsequent metamorphism at Lyagkomina.

ISOTOPIC WORK

The investigated rocks

Titanites and rutiles from the Neoarchaean rocks of the Lyagkomina Dam locality (91011 to 13 in Fig. 2) and from Tupaya Bay were separated to be investigated isotopically. Previously, zircons from the same rocks had yielded U-Pb ages in excess of 2.72 Ga (Bibikova et al. 1996).

Lyagkomina sample 91011 is a heterogeneous migmatite that has been formed from Chupa-type aluminous gneisses. It contains coarse-grained, white or slightly yellowish pegmatitoid veins and quartz-feldspar segregations. The migmatitic matrix contains small lenticular slivers of metasedimentary protolith separated by zones of quartz- and muscovite-bearing foliated rock. These zones trend parallel to the migmatitic banding and were presumably formed more or less simultaneously with the pegmatitic veins. The gneiss contains kyanite, garnet, andesine, K-feldspar, biotite and quartz with accessory zircon, apatite, titanite and rutile.

Samples 91012 and 91013 are from the central and marginal parts, respectively, of a TTG-type rock body at Lyagkomina. Sample 91012 represents a nebulitic migmatite with lenticular quartz segregations, while light-reddish feldspar bands occur in a darker bluish-grey plagioclase-rich palaeosome in sample 91013. Micas accentuate the foliation and there are also rare grains of red garnet. The rocks are migmatised tonalites to granodiorites, sometimes with preserved relics of relatively coarse-grained subhedral andesine containing fine-grained inclusions of likewise subhedral amphibole, biotite and quartz. The plagioclase is deformed and partly recrystallized. Accessory apatite and zircon ± magnetite appear to belong to the igneous association and are often enclosed within coarser-grained andesine. Rutile mostly occurs as inclusions in the quartz crystals. Titanite, biotite intergrown with muscovite, albite, and fine-grained muscovite convey a banded character to these rocks.

Apart from Neoarchaean supracrustal and plutonic lithologies, the Lyagkomina locality also features later mafic dyke intrusions. Sample 91014 is from such a dyke of approximately dioritic chemical composition that has been intruded into the local Chupa-type aluminous gneisses. In spite of remains of igneous plagioclase, the present mineral set-up is wholly metamorphic, garnet and hornblende being the most prominent dark phases, with some clinopyroxene similarly present. The metamorphic mineral association also includes most of the zircons. While there is age evidence that some zircon grains have been captured from older rocks (cf. below), virtually none of those analysed appear to have formed during the igneous crystallization of the dyke. Additional particulars of chemistry and mineral composition are given in the following sections on P-T work and U-Pb data (see also Table 1).

Compared with the strong Palaeoproterozoic re-

working at Lyagkomina, the one at Tupaya Bay has been relatively weak. In a previous study of the latter region, Bibikova et al. (1996) could date several Archaean igneous and metamorphic events using zircon U-Pb chronology. That paper and several others (e.g. Bibikova et al. 1993, Bogdanova & Bibikova 1993, Lobach-Zhuchenko et al. 1993) in addition provide accounts of the Neoarchaean and later lithologies now sampled from the Tupaya Bay area. In the present study, titanites were dated also from the ca. 2.45 Ga old potassic granites (cf. Bibikova et al. 1993) close to Tupaya Bay (Fig. 1).

Methods

The chemical preparation of the mineral samples selected for radiometric age determination followed conventional isotope dilution methods originally described by Krogh (1973 and 1982). The total Pb blank for our small-scale zircon procedure was 7 to 15 picogram during the course of this project. Additional steps involving treatment with bromic acid increased the blank level of the titanites and rutiles to 10-20 picogram. Isotopic data were obtained in static mode on samples mixed with 205-233-235 tracer using a Finnigan 261 spectrometer. Data processing was carried out according to Ludwig (1991a and b), using the decay constants recommended by Steiger and Jäger (1977).

P-T WORK ON METAMORPHIC MINERALS

The P-T conditions of the investigated mineral associations in the metamorphosed Lyagkomina amphibolitic dykes were assessed by employing the Perchuk et al. (1985) garnet-amphibole thermometer, the Kohn & Spear (1990) garnet-plagioclase-amphibole barometers, and the Plusnina (1982) amphibole-plagioclase thermobarometer. The computer program "Geothermobarometry", coded by M. Kohn (Spear et al. 1991), was also employed. Estimates of the P-T conditions for the peak and retrograde stages of metamorphism were obtained from core-core and rim-rim mineral relationships, respectively.

In contrast to the Neoarchaean, often nearly ultramafic amphibolites in the region (cf. Stepanov & Slabunov 1989), the Lyagkomina dykes are more silicic, with SiO₂-contents between 54.0 and 55.6% and Na₂O+K₂O contents between 1.84 and 3.18%. The mineral assemblages evidence a less complex metamorphic history than that of the Archaean amphibolites. The igneous assemblage is represented by relic subhedral plagioclase. Otherwise, the rocks mainly compose garnet, hornblende, rare clinopyroxene, labradorite, biotite and quartz. Zircon, allanite, apatite, opaques and rutile are accessories. Small amounts of titanite are also present. Scapolite, calcite, leucoxene and chlorites are partly replacing plagioclase, hornblende and biotite.

RESULTS

Stages and P-T conditions of metamorphism

Microprobe analyses of minerals from the garnetamphibolitic mafic dyke at Lyagkomina (sample 91014) are reported in Table 1.

Two metamorphic mineral assemblages, which were formed at somewhat different times, can be distinguished. The early one is represented by relatively coarser-grained subhedral garnet (Pyp17-19Gros32-33Alm44-46Spes3-4), pargasitic hornblende (Al_{tot}2.5-2.7), biotite, plagioclase (An49-54) and quartz (see Table 1). These minerals form coarse intergrowths and define the nematoblastic texture of the rock. This earlier assemblage yield-

Table 1. Representative microprobe analyses of metamorphic minerals from the Lyagkomina dyke.

					1	Peak ass	emblages						
Mineral*	grt 7	grt 11	grt 12	grt39	grt40	grt41	amph 22	amph23	amph25	amph63	p131	p132	p153
SiO2	37.66	38.5	37.9	38.11	38.02	38.15	43.66	43.1	43.41	43.16	55.09	55.17	53.19
A12O3	21.73	21.86	21.75	21.69	21.7	22.21	14.53	15.23	15.2	15.13	28.73	28.43	28.8
FeO	21.23	21.43	21.85	21.91	21.94	22.38	13.49	13.64	14.03	13.53	0.17	0.15	0.08
MgO	4.81	5.06	4.66	4.66	4.72	4.82	11.26	11.04	10.91	10.94	0.03	0.01	0.07
CaO	11.72	11.57	11.51	11.62	11.71	11.35	11.91	12.11	12.14	12.18	10.56	10.17	11.11
Na2O	0	0.07	0.05	0	0.08	0.04	0.99	1.09	1.12	1.07	5.55	7.72	4.92
K2O	0.02	0.01	0.03	0.01	0	0.03	0.64	1.05	0.84	1.42	0.09	0.08	0.04
TiO2	0.7	0.05	0.02	0.04	0.05	0.02	0.93	0.93	0.94	1.14	0.03	0.02	0.03
MnO	1.32	1.2	1.53	1.53	1.42	1.46	0.25	0.24	0.22	0.14	0.01	0.03	0.01
Cr2O3	0.02	0.01	0.02	0.16	0.06	0.05	0.08	0.15	0.15	0.11	0.01	0.01	0.01
Si	2.941	2.981	2.957	2.964	2.954	2.941	6.4	6.295	6.316	6.298	2.476	2.462	2.442
Al	2	1.994	2	1.988	1.987	2.018	2.51	2.622	2.606	2.601	1.522	1.495	1.558
Fe3	0.036	0.049	0.092	0.071	0.108	0.104	0	0	0	0	0	0	0
Fe2	1.351	1.339	1.334	1.355	1.317	1.339	1.654	1.666	1.707	1.651	0.006	0.006	0.003
Mg	0.56	0.584	0.541	0.54	0.546	0.553	2.461	2.404	2.367	2.379	0.002	0.001	0.005
Ca	0.981	0.96	0.962	0.968	0.975	0.937	1.87	1.895	1.892	1.903	0.509	0.486	0.547
Na	0	0.01	0.007	0.001	0.012	0.006	0.281	0.309	0.315	0.304	0.484	0.668	0.438
Κ	0.002	0.001	0.003	0.001	0	0.003	0.12	0.196	0.156	0.264	0.005	0.004	0.002
Ti	0.041	0.003	0.001	0.002	0.003	0.001	0.102	0.102	0.103	0.125	0.001	0.001	0.001
Mn	0.087	0.078	0.101	0.101	0.094	0.095	0.032	0.03	0.027	0.017	0	0.001	0.001
Cr	0.001	0.001	0.001	0.01	0.004	0.003	0.01	0.017	0.017	0.012	0	0	0
mg	29	30	29	29	29	29	60	59	58	59	25	15	61
Albite	•	•	•	•	•	•	•	•	•	•	48.5	57.7	44.4
Anorthite	•	•	•	•	•	•	•	•	•	•	51.0	42.0	55.4
Orthoclase	•	•	•	•	•	•	•	•	•	•	0.5	0.4	0.2

ed a P-T estimate of 650±30°C at 8.5-9.5 kbar.

The apparently later metamorphic assemblage is made up of light-coloured hornblende, which is more edenitic and contains less Ti than the early one, garnet (Pyp16-18Gros27-30Alm48-49Spes4-6), labradorite (An60-66) and quartz. These minerals form intergrowths that surround and separate early garnet from hornblende and biotite, a situation we believe to be related to a decompressional trend in the metamorphic evolution. This particular stage is estimated to have occurred at $630\pm20^{\circ}$ C and 7-8 kbar.

The obtained difference between the two metamorphic stages, which represent roughly similar temperatures at falling pressures, indicates nearisothermal decompression and rapid uplift in a collisional-type environment. The presently recorded metamorphic conditions in the dyke are



Fig. 3. Concordia diagram for zircons of sample 91014 from the Lyagkomina area. The discordia is based on the four brilliant-type zircon analyses reported in Table 2.

* grt – garnet, amph – amphibole, pl – plagioclase. All mineral calculations are made by MINTAB software. The minerals were analysed with GEOL-JSM-6400 (Lund University) using natural standards and ZAF correction program.

				Retrogra	ade assemb	olages				
grt13	grt14	grt43	grt61	grt70	amph47	amph 48	amph68	pl15	pl17	p152
37.9	37.82	38.48	38.26	38.36	41.73	42.23	42.45	52.19	51.8	50.78
21.69	21.57	22.08	21.92	22.12	15.45	15.13	15.27	31.34	31.08	31.02
21.69	22.37	21.75	22.39	21.79	14.31	13.84	13.77	0.24	0.17	0.09
4.58	4.58	4.99	4.17	4.73	10.41	10.48	10.83	0.01	0.17	0.01
12.02	10.89	12.13	11.82	12.1	11.76	11.91	12.13	12.39	13.09	13.3
0.49	0.08	0.06	0.02	0.05	1.22	1.17	1.15	4.04	4.03	3.79
0.01	0.01	0.04	0.01	0	1.09	0.88	1.39	0.53	0.04	0.03
0.07	0.1	0.02	0.04	0.02	0.94	0.9	1.07	0.56	0.01	0.01
1.48	1.87	1.26	2.01	1.41	0.26	0.25	0.2	0.02	0.03	0.07
0.13	0.2	0.08	0.2	0.05	0.17	0.18	0.15	0.02	0.02	0.03
2.925	2.954	2.948	2.964	2.95	6.209	6.281	6.238	2.34	2.342	2.327
1.973	1.986	1.993	2.002	2.005	2.71	2.652	2.645	1.656	1.656	1.676
0.236	0.094	0.116	0.098	0.098	0	0	0	0	0	0
1.164	1.367	1.278	1.451	1.304	1.781	1.722	1.693	0.009	0.006	0.003
0.527	0.533	0.569	0.481	0.543	2.309	2.323	2.372	0.001	0.012	0.001
0.993	0.912	0.995	0.981	0.997	1.875	1.898	1.909	0.595	0.634	0.653
0.073	0.012	0.009	0.003	0.008	0.352	0.338	0.329	0.351	0.353	0.337
0.001	0.001	0.004	0.002	0	0.207	0.167	0.261	0.03	0.002	0.002
0.004	0.006	0.001	0.002	0.001	0.106	0.1	0.118	0.019	0	0
0.096	0.124	0.082	0.132	0.092	0.033	0.032	0.025	0.001	0.001	0.003
0.008	0.012	0.005	0.010	0.003	0.02	0.021	0.017	0.001	0.001	0.001
31	28	31	25	29	56	57	58	10	65	19
•	•	•	•	•	•	•	•	36.0	35.7	34.0
•	•	•	•	•	•	•	•	60.9	64.0	65.8
•	•	•	•	•	•	•	•	3.1	0.2	0.2

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	(a)	and rock	(mg)	(p) (b)	(q) (mdd)	(mol%) rad.(c)	measured	(p)	±2s%	(p)	±2s%	(p)	±2s
98031 ti	y, frag	Tup, K-gr	2.00	22	∞	12.0	256	0.3338	0.30	5.204	0.50	1849	2
98029 ti	y, frag	Tup, K-gr	2.13	18	9	10.0	218	0.3327	0.30	5.174	0.50	1845	4
96116 ti	y, +100, frag	Tup, diorite	0.84	14	4	6.3	105	0.2990	0.59	4.672	0.60	1855	6
96116 ti	y, -100, frag	Tup, diorite	0.81	23	8	8.2	255	0.3301	0.45	5.119	0.49	1840	З
91013 ti	bry, frag	Lyag, ton	0.52	82	31	13.4	750	0.3359	0.53	5.292	1.04	1868	15
91011 ru	y, honey, nee	Lyag, Algn	0.28	37	11	3.4	1700	0.3136	0.29	4.650	0.53	1758	~
91012 ru	ybr, nee,-100	Lyag, ton	0.63	15	2	1.6	530	0.3173	0.27	4.718	0.62	1763	6
91012 ru	frag, tab +100	Lyag, ton	1.06	18	9	3.0	1150	0.3158	0.24	4.737	0.46	1779	2
91014 ru	ybr, nee	Lyag, dyke	0.26	16	5	2.0	440	0.3216	0.49	4.750	0.86	1751	12
95010 ru	gr,	Tup, gabbro	1.69	10	З	0.7	1775	0.3099	0.43	4.551	0.70	1740	6
92077 ru	red	Tup, Algn	1.76	60	18	0.1	11000	0.3149	0.40	4.674	0.60	1760	С
95005 ru	red	Tup, Algn	1.84	38	11	1.0	2920	0.3124	0.40	4.606	0.50	1748	0
95005 ru	gr,	Tup, Algn	1.52	14	13	1.0	6500	0.3131	0.50	4.618	0.70	1749	2
91014 z, M	brill, $ab +60$	Lyag, dyke	0.02	186	60	0.1	4000	0.3371	0.87	5.329	0.94	1874	9
91014 z, A	brill, ab –74	Lyag, dyke	0.12	62	20	1.4	380	0.3312	0.34	5.257	1.12	1882	17
91014 z, E	brill, ab+90, 15xx	Lyag, dyke	0.07	72	21	0.6	740	0.3087	0.49	4.850	1.05	1863	15
91014 z, H	brill-60	Lyag, dyke	0.11	93	25	1.3	486	0.2725	0.31	4.276	1.15	1861	18
91014 z, D	pi, frag, core	Lyag, dyke	0.07	268	<i>4</i>	3.8	570	0.2927	0.22	5.026	0.88	2022	13
91014 z, L	lp, +90, 9xx	Lyag, dyke	0.02	256	95	1.1	1570	0.3750	0.62	7.248	0.65	2230	З
91014 z, C	lp, +60, 33xx	Lyag, dyke	0.09	189	57	0.4	1480	0.3124	0.19	4.904	0.41	1862	9
91014 z, K	lp, -60, 62xx	Lyag, dyke	0.06	256	75	1.1	1910	0.3023	0.21	4.996	0.30	1954	З
(a) all zircon	crystals are transpe	arent. uncolour	ed and no	nmagnet	ic (zero s	lone. 3 deør	of tilt and 1.7	A of Frantz is	odvnamic	senarator) unle	ess otherv	vise indica	ted:

zzircon; ti=titanite; ru=rutile; Capital letter=separate fraction; +60=size in microns; 11xx=number of crystals analysed; br=brown; gr=green; y=yellow; pi=pink; nee=needle; lp=long-prismatic(l/w≈4 or more); brill=brilliant roundish; frag=fragment; tab=table-like; ab=abraded (Krogh 1982); core=cores or inner structures are visible; K-gr=potassium granite; ton=tonalite; dyke= amphibolitic dyke Algn=aluminous gneiss

⁽b) Concentrations are known to about 5% for sample weights of about 0.1 mg, and to about 10% for sample weights of about 0.05 mg

⁽c) Relative to total radiogenic Pb, and a measure of approximate Th content and U/Th ratio (d) Corrected for lead blank of 5-15 pg, fractionation, spike and initial common lead (Stacey & Kramers 1975); errors are at the 2 sigma or 95% level and are expressed in per cent of the atomic ratio and in absolute numbers for the age. All rocks but 91014 and the 2450 Ma K-granites are of Archaean origin.



Fig. 4. Transmitted light photographs of the type of zircons and rutiles present in the rock samples. The centralupper part exhibits some rejected and not analysed zircon crystals (ca. 90µm wide) with obvious cores. Abbreviations: lp=long prismatic; brill=brilliant type as in Table 2.

similar to those previously assessed for the latestage metamorphism which altered the aluminous kyanite-biotite-muscovite Chupa gneisses at Lyagkomina (Bibikova et al. 1993).

U-Pb isotopic data

All the U-Pb isotopic data and ages are presented in Table 2, and those for the zircons are plotted in a Concordia diagram in Figure 3. Figures 4 and 5 display photographs of minerals of the kind analysed, like some of those rejected.

Zircons.

The zircon crystals from the investigated mafic dyke show only limited variability in morphology and style of crystallization. Most of them are transparent, short prismatic to isometric and have well-developed euhedral shapes, often with a high



Fig. 5. Transmitted light photographs of the type of titanites analysed. Magnification as in Figure 4. Abbreviations correspond to those in Table 2.

lustre (lower left in Fig. 4). Clearly, these are not common features among mafic-dyke zircons. A minor proportion of the grains is elongated, longprismatic (upper left and centre of Fig. 4).

Some of the zircon analyses divert towards higher Pb/Pb ages (> 1900 Ma) than those of the transparent crystals, which in turn delimit the concordia-discordia upper-intercept age to 1875 ± 4 Ma (at 95% confidence level and with a MSWD of 1.3).

Titanites.

In the present study, we have investigated titan-

ite crystals both from the Lyagkomina and Tupaya Bay localities. The titanites have been separated from rocks which according to previous studies (Bibikova et al. 1996) and zircon ion-probe data (Claesson et al. in preparation) are Archaean and early Palaeoproterozoic in age. The analytical results, together with calculated ²⁰⁷Pb/²⁰⁶Pb ages, are presented in Table 2. Like titanites in general, the presently studied crystals (some examples are shown in Fig. 5) have low concentrations of uranium, leading to highly non-radiogenic Pb-isotopic compositions. Still, all the titanites have concordant to almost concordant compositions indicating ages close to 1860 Ma.

Rutiles.

Rutile isotopic compositions and ages have been studied from some of the Archaean rocks in the study area. To judge from the high ²⁰⁶Pb/²⁰⁴Pb isotopic parameter, Pb in the rutiles is relatively radiogenic. Uranium contents are low, however, and it was a surprise to encounter concordant ages at about 1750 Ma.

Comparison with previous titanite and rutile U-Pb isotopic ages.

The titanite and rutile U-Pb ages of this study compare well with the corresponding ages obtained during the regional study of the Belomorian Belt by Bibikova et al. (2001). In that paper, the titanite ages from the central part of the belt range between 1.82 and 1.87 Ga, while our ages have a range from 1.84 to 1.87 Ga. For rutile the equivalent ranges are 1.75–1.81 and 1.74–1.78 Ga, respectively. In the present data set we find no systematic correlation between rutile ages and rock lithologies or other rock characteristics.

DISCUSSION

Evidence from the P-T study

The obtained data constrain the timing of the P-T and tectonic evolution of the crust in the Lyagkomina area. We suggest that the 1875-Ma zircon



Fig. 6. P-T-t diagram showing the metamorphic evolution of the Lyagkomina garnet-amphibolitic dyke (black arrow) in comparison with the Palaeoproterozoic P-T trend of the host Archaean garnet-kyanite-biotite-muscovite gneisses (grey arrow). The dashed arrow shows a possible further evolution of the crust until rutile in the gneisses reached its closure temperature. The striped field indicates the P-T conditions of dehydration melting of amphibolite (after Wyllie & Wolf 1993). The curves are. 1. dry gabbro solidus (Wyllie et al. 1981); 2. H₂O-saturated granite solidus (Huang & Wyllie 1973); Muscovite-out dehydration melting of muscovite (after Thompson 1990); Biotite-out dehydration melting of biotite (Patino Douce & Beard 1995).

age dates peak metamorphism in the mafic dyke. This took place close to 650°C at ca. 9 kbar (Fig. 6), but with the possibility that evidence of a still slightly higher peak temperature had been lost by element diffusion from the relatively small garnets of the dyke rock.

The indicated peak P-T conditions correspond to the earliest phase of recumbent folding of the dyke and its host rocks in the lower crust at depths of ca. 35 km. Since the peak-metamorphic mineral assemblage is well within the solidus field, igneous crystallization of the dyke-rock melt had apparently been completed at that time. During that stage, new metamorphic zircon could grow rather easily due to the decomposition of pyroxene and other magmatic host minerals of zirconium (see papers in Chemical Geology 117, 1994).

Rims of plagioclase around garnets, changes of mineral compositions and the appearance of anatectic veining in the dyke and its surroundings, all indicate sudden decompression of the rocks and jump uplift of the crust with unroofing of approximately 9-10 km during the first stage of retrogression from peak conditions. This was an almost adiabatic process with little cooling. The average titanite age of ca. 1860 Ma, with the assumption of a 650-700°C closure temperature of its U-Pb isotopic system, may indicate that this initial stage of the tectonothermal evolution lasted ca. 15 Ma or even less. This corresponds to uplift rates of ca. 0.75 mm/year which appear to be a result of tectonic erosion and/or collapse of the crust after its thickening during the Lapland-Kola orogenic collision (cf. e.g. Harley 1992).

The change from compressional to extensional regime is supported by later refolding into open folds and penetrative ductile shearing. This latter structural development and cooling of the crust can be reconstructed using P-T estimates for the aluminous gneisses which host the analysed dyke (Bibikova et al. 1993). They document a prolonged and slower tectonothermal evolution during a 110-Ma period that is constrained by the titanite and rutile ages (Fig. 6). Such situations are typical for the uplift of lower crust in most Proterozoic high-grade terrains and imply thermal relaxation of the entire lithosphere to a steady-state geothermal regime (cf. Harley 1992).

The equivalent 1.9–1.75 Ga crustal development in the Tupaya Bay locality appears to have occurred at a shallower level in the crust. Very probably this difference was controlled by the Neoarchaean prehistory of the Lake Kovdozero region when the Lyagkomina rocks were buried ca. 7 km deeper in the crust than those in the Tupaya Bay area (cf. Bibikova et al. 1993, Lobach-Zhuchenko et al. 1993).

Formation and age of the Lyagkomina maficdyke zircons

Mafic magmas are low in silica, which commonly prevents the igneous crystallization of zircon. Even where there is free SiO_2 , zirconium (Zr) is commonly partitioned into pyroxenes and other dark minerals. In the presently investigated dyke, however, zircons are abundant. They are transparent, highly lustrous, and euhedral with only a limited number of simple crystal facets. This suggests crystallization during metamorphism rather than from melt.

During igneous crystallization of mafic rocks, Zr mostly enters dark minerals, mainly pyroxenes (Poldervaart, 1956) but also oxide opaques. If, however, some Zr remains and free SiO_2 is enriched in the residual melt, zircon forms late, mostly as irregular and anhedral grains in the interstitial spaces. These are very different from the euhedral zircons found in the Lyagkomina dyke.

During metamorphism and the attendant decomposition of igneous minerals, Zr is released from pyroxene and opaques, while more SiO_2 can rather easily be introduced by circulating fluids. The observed low contents of U in the mafic-dyke zircons and their highly lustrous appearances indicate a metamorphic origin at relatively high P-T conditions (Bibikova 1984). The 1875±4 Ma age determined from the high-quality zircons (Fig. 3) is therefore taken to date an early, high-T stage.

The remaining, considerably less abundant, Uricher mafic-dyke zircons have long-prismatic shapes and yield higher 207Pb/206Pb ages of 1954 and 2022 Ma. Most probably, these grains include xenocrysts which had been absorbed into the melt from the country-rocks. Zircons of that appearance are common in the nearby Neoarchaean tonalitic gneisses where complex growth and/or Pb-loss processes appear to have occurred. Assimilation of zircon xenocrysts into various rock melts (e.g. Pidgeon & Aftalion 1978, Pidgeon et al. 1996) has been experimentally confirmed to reflect their high melting temperatures (Watson & Harrison 1983). These are reasons why zircon xenocrysts can survive long in Zr-saturated melts. They are common in granitic rocks and can occasionally complicate multi-grain U-Pb dating (cf. Eliasson & Schöberg

1991). Felsic rocks are of course most easily saturated in Zr, but xenocrysts have even been described from rapidly crystallized komatiites (Compston et al. 1985).

Experimental studies on the preservation of zircon U-Pb isotopic systems during melting (e.g. Bibikova et al. 1995) indicate that, although radiogenic Pb is easily lost from metamict domains, textures of crystalline zircon often remain unaffected even during prolonged heating to temperatures in excess of 1000°C. It is plausible, however, that long-lasting elevated P-T conditions cause resorption of zircon xenocrysts if fluids are present, and that there is crystallization of new overgrowths on zircon seeds or entirely new crystals are formed (Skiöld 1986). We therefore consider that the presently analysed zircon xenocrysts may have preserved an isotopic memory of their primary crystallization.

Cooling history as reflected by the ages of zircon, titanite and rutile

Titanite growth and recrystallization can provide age estimates for deformational events associated with tectonometamorphic processes (see Resor et al. 1996, Getty & Gromet 1992). Similarly, titanite and less commonly rutile have been used in geochronological studies for the determination of cooling histories in metamorphic terrains. In early studies (e.g. Mattison 1978, Tucker et al. 1987), the closure temperatures of U and Pb isotopes in titanite were considered to range between ca. 550 and 600°C. Later petrological-geochronological experiments, however, indicated higher temperatures of approximately 650°C (Mezger et al. 1991) and in excess of 700°C (Scott & StOnge 1995, Zhang & Schärer 1996, Pidgeon et al. 1996).

Diffusion parameters for Pb in variously-sized titanites have been determined by Cherniak (1993) using the diffusion equation of Dodson (1973). Cherniak's calculations for 300- μ m radius crystalline titanite in a rock with a cooling rate of 2°C/ Ma suggest a closing temperature just above 700°C. In comparison, the closure temperature for rutile is considerably lower. According to Mezger et al. (1989) U-Pb diffusion from 90-200 μ m rutiles with a cooling rate of 1°C/Ma closes at about 420°C.

From petrographic investigation we conclude that the rutiles and titanites as well as the highly lustrous dyke zircons from Lyagkomina are all metamorphic. Although the titanites from both Lyagkomina and Tupaya Bay are low in uranium, which causes relatively large error, the data obtained are mostly concordant at approximately 1860 Ma. This means that ca. 1860 Ma ago the metamorphic temperature in the Lyagkomina-Tupaya Bay area passed titanite blocking conditions at approximately 700°C. The rutile age estimates are concordant at about 1750±10 Ma.

These observations and age determinations allow the estimation of cooling rates after peak metamorphism. The P-T measurements on mineral pairs indicate a metamorphic peak close to $650 \pm$ 30° C and ca. 9 kbar with possibly still higher maximum temperatures. This compares well with the experimental data preferring a closing temperature of roughly 700°C for titanite. Assuming a constant rate of cooling over the temperature interval from 700-650 to 400°C between 1860 and 1750 Ma, a ratio of approximately 2.5°C/Ma is obtained. However, cooling rates are more likely to vary over such a wide range of temperatures, being higher at the elevated temperatures that prevail during the early stages of cooling.

The difference between the closure ages for metamorphic zircon and titanite is not great but quite obvious. The mafic-dyke zircons of the Lyagkomina area are ca. 15 Ma older than the metamorphic titanite ages from the 2.8-2.4 Ga country rocks. Nevertheless, it is unlikely that this age difference can be used to model cooling rates, where e.g. cooling from 850 to 700°C in 15 Ma would suggest ca. 10°C/Ma. The reason is that we cannot anticipate or exclude that the highly lustrous zircons had grown after passing their blocking temperature in the environment prevailing in the studied area. In the discussion above we rather suggest that they crystallized from Zr released during metamorphism from its original igneous host minerals. Circulating metamorphic fluids have most probably contributed in such a process.

SYNOPSIS AND CONCLUSIONS

Combined U-Pb isotopic and P-T studies of minerals and mineral associations in the northern part of the Belomorian Belt have succeeded to assess the metamorphic evolution of that region in the late Palaeoproterozoic, in particular the time which followed upon the stacking and thickening of the Belomorian crust during the Lapland-Kola continental collision. The most important conclusions of the present study are:

- Highly lustrous metamorphic zircons in a Palaeoproterozoic mafic dyke now recrystallized into garnet-amphibolite reflect a stage of late Palaeoproterozoic metamorphism which peaked at approximately 1875±4 Ma ago.
- Peak metamorphic conditions were in excess of ca. 650°C at 8.5 to 9.5 kbar, whereas mineral associations formed somewhat later indicate a following stage at ca. 630°C and 7-8 kbar. Metamorphism thus occurred at a maximum depth of ca. 35 km.
- In conjunction with the new isotopic ages, the obtained P-T data indicates a relatively shortlived stage of initially rapid, almost adiabatic decompression. This suggests sudden jump uplift of previously thickened crust and unroofing of ca. 9–10 km.
- Cooling immediately after the initial stage of decompression and strong crustal uplift were rapid processes. The rate then slowed down substantially as indicated by titanite U-Pb ages of ca. 1860 Ma and rutile ages of ca. 1750 Ma. During the ca. 110-Ma time interval between the indicated ages, the average cooling rate was ca. 2.5°C/Ma, presumably higher in the beginning of the time interval and lower at the end.
- The titanite and rutile U-Pb ages measured in the present study virtually coincide with the ages previously obtained in the central Belomorian Belt during a much wider regional survey (Bibikova et al. 2001).
- Differences in titanite and rutile U-Pb ages as a function of lithological variations seem to be absent.

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